Certificate

Quality Management System
EN ISO 13485:2016
EN ISO 13485:2016/AC:2018
EN ISO 13485:2016/A11:2021

Registration No.:	SX 2468805-1
Certificate Holder:	Cretex Medical Component and Device Technologies, Inc. d.b.a. JunoPacific, Inc. 8701 95th Avenue North Brooklyn Park MN 55445 USA
Scope:	Contract Manufacturing of sterile and non-sterile custom OEM injection molded components and assemblies for Medical Devices under environmentally controlled conditions. Contract Manufacturing and Distribution of Sterile, Non-active Finished Medical Devices.

The Certification Body of TÜV Rheinland LGA Products GmbH certifies that the organization has established and applies a quality management system for medical devices.
Proof has been furnished that the requirements specified in the abovementioned standard are fulfilled. The quality management system is subject to yearly surveillance.

Report No.:	234220380-11
Effective date:	2025-02-24
Expiry date:	2028-02-23
Issue date:	2025-02-04

Replaces certificate SX 2468805-1 issued 2024-05-10.

This certificate can be validated on https://www.certipedia.com

Melissa Di Buono-Russo
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

DAkkS Deutsche Akkreditierungsstelle D-ZM-14169-01-02

TÜVRheinland®
Precisely Right.